







# Lunar Polar Volatiles: Assessment of the Current State of Knowledge

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### Introduction

# Background

- > Lunar volatiles are potentially a valuable resource for exploration.
- Understanding of volatiles is evolving as new data emerge and scientists synthesize data, theory, lab experiments, and models.
- Data sets are nuanced in ways that are not always obvious to the outside observer.

# Purpose

- To provide an overview of the present state of knowledge
- To explain the level of certainty/uniqueness of the results
- > To interpret the findings for application to ISRU

# Volatiles on the Moon

### Three brands of volatiles:

- > Sequestered volatiles in cold traps
  - Episodic delivery of large quantities or constant delivery of small quantities or both
- Internal volatiles trapped in minerals and glasses
  - Leftover from lunar formation
- > Global surface volatiles
  - Transient veneer either produced and lost in place diurnally or involved in migration



# Measurements of Polar Volatiles (Water Ice)

- Current focus: Obtaining measurements of the following quantities
  - > Abundance
    - Exploration rationale: assess value and extraction technique
    - Science rationale: tied to abundance of sources
  - > Composition
    - Exploration: additional resources
    - Science: chemical fingerprint of source
  - > Distribution
    - Exploration: scale of operations and extraction technique
    - Science: age of deposits; redistribution and retention processes
  - > Physical State
    - Exploration: extraction technique; operational paradigm
    - Science: interactions between volatiles and regolith

# Processes for Polar Volatiles (Water Ice)

 Current focus: Understanding the lunar water system from source to sink

Abundance

Composition

Distribution

Physical Form

SOURCES

Comets
Meteoroids
Outgassing/internal
Solar wind

PRESENT DAY LUNAR VOLATILES

REDISTRIBUTION

Migration/hopping
Migration/thermal vertical
Impact gardening

**SINKS** 

Sublimation/thermal desorption Photodissociation

Impact

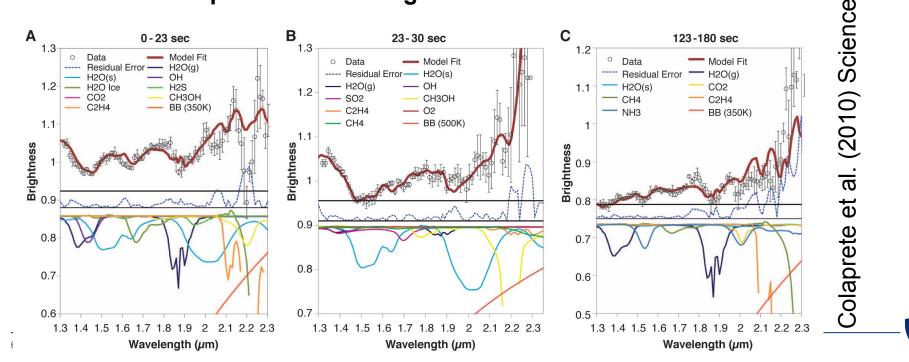
Sputtering

Chemistry

# Existing Data: COMPOSITION

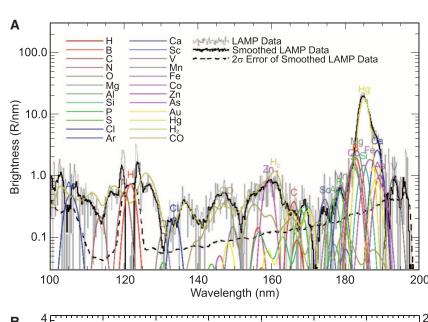
- LCROSS impact into Cabeus
- Spectral identification of water in both solid and gas phases
  - High resolution spectra provide strong evidence for the chemical composition
  - LCROSS measurement has advantage of having lofted material into sunlight providing a strong illumination source

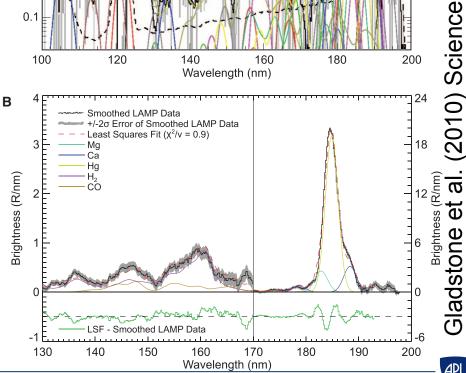
Question arises about any impact-induced chemistry and contributions from the impactor vs. the target material.



# Existing Data: COMPOSITION

- LCROSS impact into Cabeus
- Spectral identification of other volatiles in the vapor
- From LCROSS
  - > H<sub>2</sub>S, SO<sub>2</sub>
  - > NH<sub>3</sub>, CO<sub>2</sub>
  - > Hydrocarbons
  - > OH
- From LRO
  - > Hg, Mg, and/or Ca
  - > CO
  - $\succ H_2$
- From ground
  - > Na

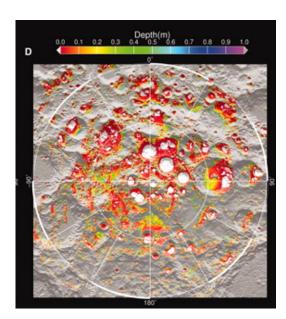




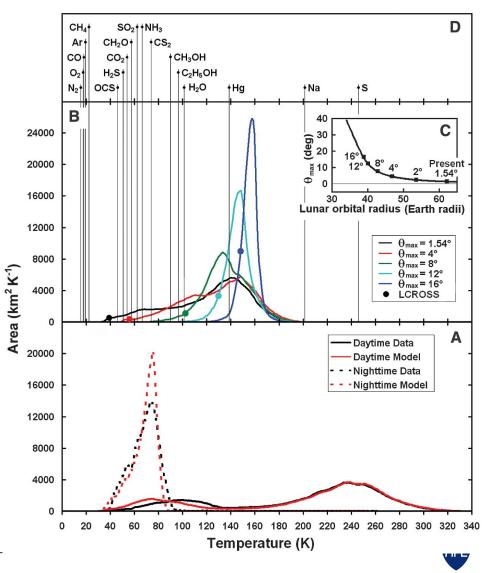
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# Existing Data: COMPOSITION & DISTRIBUTION

- LRO Diviner temperature measurements and thermal analysis
  - Provides map of where certain compounds are stable against sublimation
  - Modeling suggests the depth to a thermally stable layer

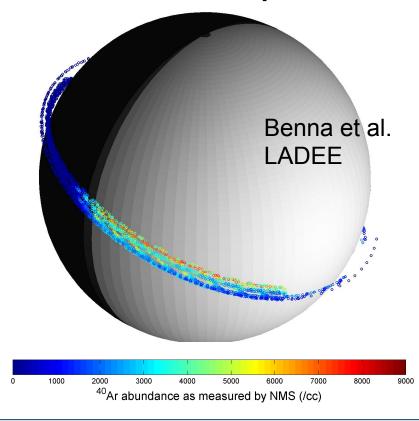


Paige et al. (2010) Science

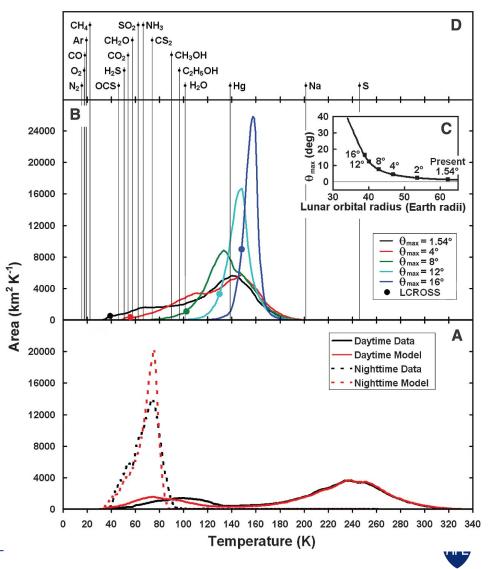


# Existing Data: COMPOSITION & DISTRIBUTION

- However we know that some exospheric species adhere to the surface at higher temperatures
- Timescale and surface interactions are important



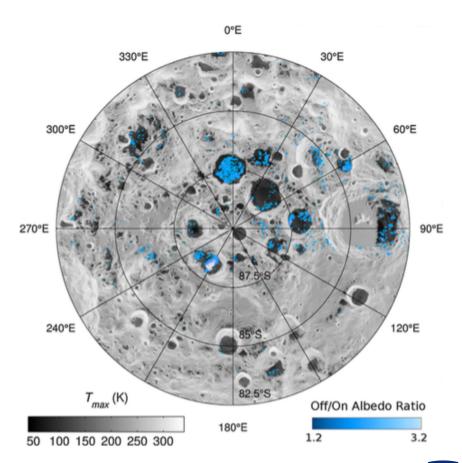
### Paige et al. (2010) Science



# Existing Data: DISTRIBUTION

- Heterogeneous lateral distribution
  - > Within a single PSR
  - > From one PSR to another
- Surface frost
  - Using the signature of water ice at 165 nm, this analysis shows that surface frost is not evenly distributed in cold regions
  - LRO LAMP data are low illumination with very coarse spectral binning, thus are better as supporting data than standing alone

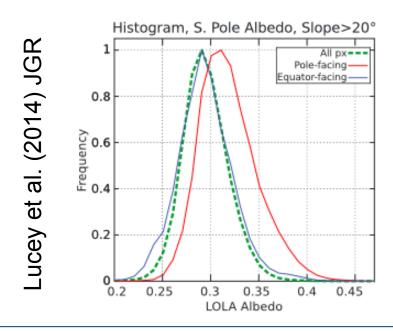
Hayne et al. (2015) Icarus

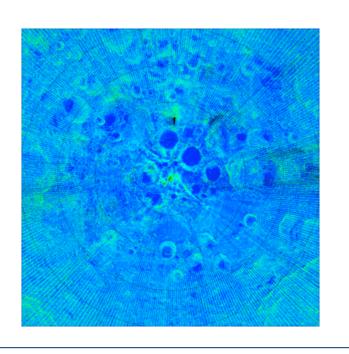


# Gladstone et al. (2012) JGR

# Existing Data: DISTRIBUTION

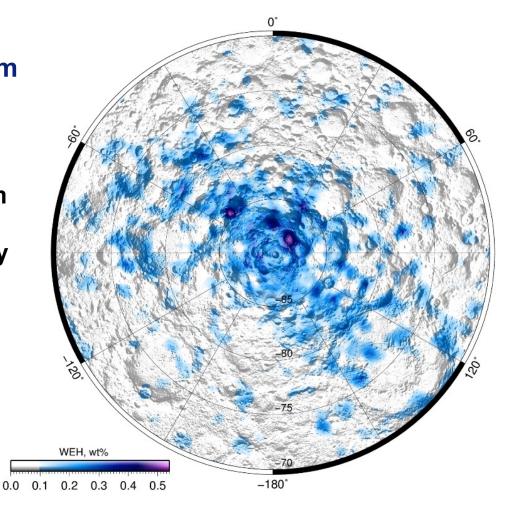
- Apparent difference in equator facing slopes and poleward facing slopes
- LRO LAMP Lyman Alpha albedo and LOLA 1064 nm albedo show difference in poleward facing slopes
- Multiple effects can produce these including the presence of frost, but no certain conclusions can be drawn on these data alone.





# Existing Data: DISTRIBUTION

- Neutron spectroscopy senses through the top 1 m of regolith.
  - Hydrogen values from depth-integrated measurements differ from surface frost measurements, especially in Shoemaker, Haworth, and Cabeus.

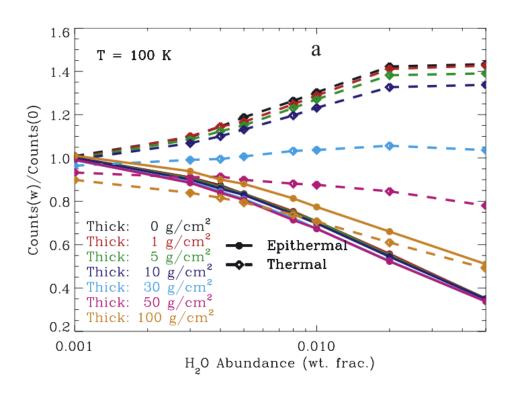


Sanin et al., 2016



# Existing Data: DISTRIBUTION

- Neutron spectroscopy senses through the top 1 m of regolith.
  - Comparing neutron data from different energy ranges gives idea of depth distribution.
  - Many sites are most consistent with a dry layer about 10 cm thick over top of a layer with higher hydrogen abundances.



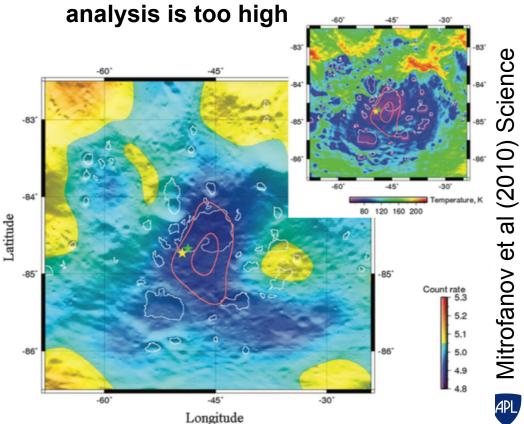
Lawrence et al. (2006) JGR

# Existing Data: DISTRIBUTION/ABUNDANCE

- Sub-pixel heterogeneity
- The LCROSS plume was consistent with 5% ± 3% water
  - Impact excavated to a depth of ~3 m and diameter ~20 m
- LEND neutron data are consistent with 0.45% water in Cabeus
  - Neutrons are sensitive to hydrogen content in top m
  - Neutron spatial resolution is > 20 m

- Possible explanations
  - Impact site was enriched laterally compared to surroundings
  - > Water is enriched below 1 m depth

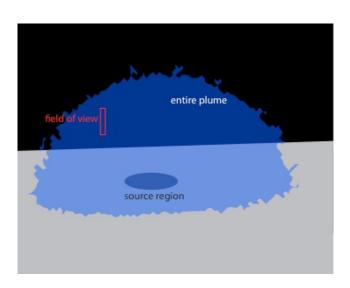
> Inferred abundance from LCROSS



# From Observations to Abundances

### **Observation of released vapor from LCROSS**





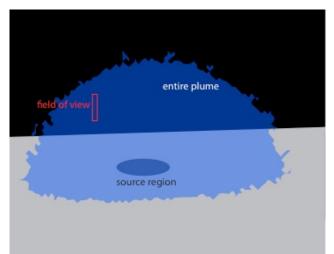
Determine volatile abundances in the regolith

# From Observations to Abundances

### **Observation of released vapor from LCROSS**

### Scalings:

Scale small field of view to global release Relate to volume of regolith affected Estimate efficiency of release



# Determine volatile abundances in the regol

LCROSS estimate of 5% water may not account for all of these

et al.	JGR
Hurley	(2012)

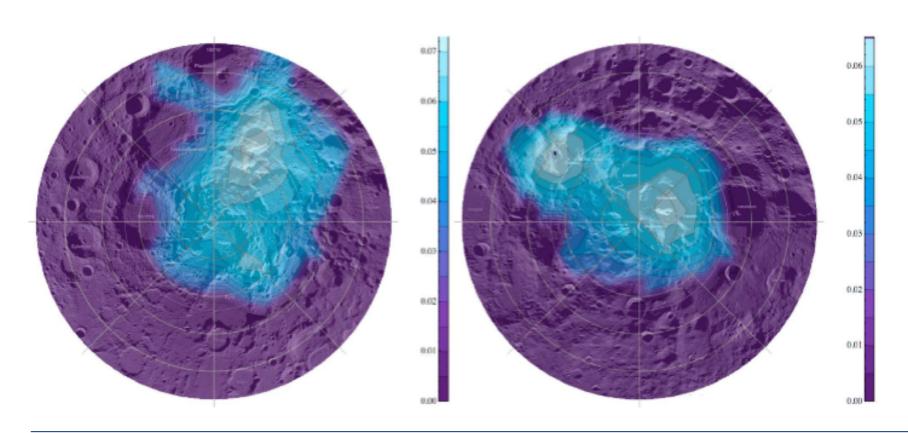
Species	Predicted amount (wt.%)	Mass (kg) released	Mass of regolith (kg)
Ca	11 <sup>C</sup>	16	140
Mg	3.4 <sup>C</sup>	3.8	110
Hg	0.28 <sup>D</sup>	12.4	4400
$H_2$	0.047 <sup>A</sup>	117	2.5e5
CO	0.023 <sup>B</sup>	41	1.8e5

# Miller et al (2012) JGR

# Existing Data: ABUNDANCE

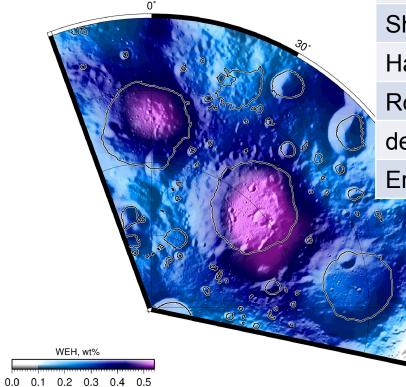
### Neutron data

- Average 0.01% wt. water equivalent hydrogen poleward of 80° and in the top 1 m
- > If water, amounts to 9.8 x 10<sup>10</sup> kg (1/1000 of Lake Tahoe or .3 mErie)



# **Existing Data: ABUNDANCE**

Crater Name	Mass of water in top 1 m (kg)	μEries (10 <sup>-6</sup> Lake Erie)	Lincoln Memorial Reflecting Pools
Cabeus	$7.7 \times 10^{10}$	160	3900
Amundsen	$5.1 \times 10^{10}$	110	2600
Shoemaker	$1.9 \times 10^{10}$	40	970
Haworth	1.6 x 10 <sup>10</sup>	34	820
Rozhd. U	$1.0 \times 10^{10}$	21	510
deGerlache	5 x 10 <sup>9</sup>	11	270
Erlanger	7 x 10 <sup>8</sup>	1.5	37



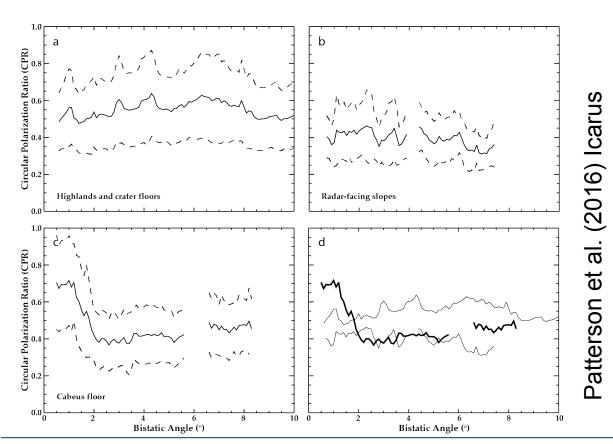
Sanin et al (2016) Icarus

# Existing Data: PHYSICAL FORM

- Radar data
- Coherent backscatter is sensitive to relatively pure blocks of ice of scale > 10 cm.
  - > On Mercury, radar data are consistent with thick, continuous, pervasive ice sheets in cold regions.
  - On the Moon, some craters show an anomalous signal where high circular polarization ratio (CPR) is observed inside the crater but not in the ejecta, unlike the majority of fresh craters
    - Spudis et al. interpret these as craters that have ice at the bottom.
    - This a controversial interpretation.
- We can rule out pervasive "skating rinks" on the Moon.

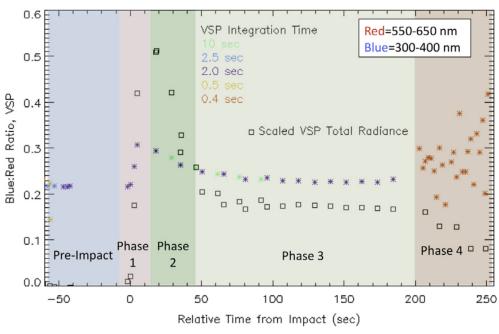
# Existing Data: PHYSICAL FORM

- Bistatic radar uses the change in circular polarization as a function of phase angle to distinguish rock from ice.
  - > Mini-RF data from Cabeus are consistent with ice present on the floor of Cabeus.



# Existing Data: PHYSICAL FORM

- LCROSS Visual Spectrometer
- Increasing Blue/Red interpreted as decreasing grain size from sublimating ice
- At least some ice is present in grains.



### **Basic Questions**

# • What is the composition?

### **Answer**

- Water ice confirmed via LCROSS & LADEE
- Hydrogen concentrations exist
- Hydrogen bearing material may or may not be water ice
- Other constituents include Hg, CO, H<sub>2</sub>S, NH<sub>3</sub>, and potentially some hydrocarbons

- More complete identification of molecular and isotopic constituents and variations with position and time
- In situ sampling, sample return, isotopic analysis, active spectroscopy

### **Basic Questions**

What is the present-day distribution?

### **Answer**

- Heterogeneities exist on many scales
  - > Latitudinal
  - Orientation of slope
  - From one PSR to another
  - > Within a PSR
  - On lateral scales of < 1 km</li>
  - As a function of depth

- Higher spatial resolution data on volatiles on the surface and with depth in lunar polar regions
- In situ sampling with subsurface access, radar, higher spatial resolution mapping



### **Basic Questions**

What is present-day abundance?

### **Answer**

- Poleward of 80°,
   ~10¹¹¹ kg of water
- Within PSRs,1-2% by weight ifH is in form ofwater
- Surface measurements consistent with <</li>2% frost

- Understanding of the chemical form of H and the overall distribution
- In situ sampling, radar, remote sensing

### **Basic Questions**

# • What is the physical form?

### **Answer**

- Pervasive, coherent ice sheets have been ruled out
- Some smaller ice grains have been detected
- Small amounts of surface frost are possible
- Pore-filling ice and hydrated minerals are also possible

- Ground truth;Laboratoryanalysis ofanalogs
- In situ sampling, sample return, active spectroscopy

# Volatiles: Future Prospects

- Resource Prospector to provide in situ "ground truth" on composition, distribution, abundance, physical form
  - > Much anticipated!
  - > Although sampling in permanent shadow is a goal, it will not do much exploration of them, or even sample the coldest regions

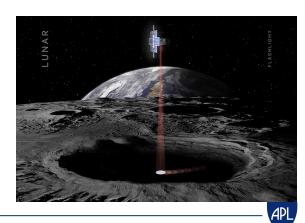
### Cubesats

- Providing more information of surface veneer, surface frost, and hydrogen
- > Will improve understanding of distribution and composition









# Volatiles: Future Prospects

- Additional, dedicated lunar volatiles missions are needed
  - > Orbital
  - > Landed
  - > With exploration and science objectives
- Leveraging multiple techniques is key
  - > Want information on surface content and volume content
  - > Spectral data and in situ identification are complementary
- Understanding the system and processes is important
  - > Follow up on new information on sources
  - > Observe the redistribution process
  - Loss processes are linked to present distribution

